

TITLE: Purification and properties of L-
asparaginase
produced by Aspergillus niger, S-48
TAT, the causal fungus of
biodeterioration inside Tut
Ankhamen Tomb (TAT)
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AB The purification and properties of L-asparaginase (I)
produced by A.
niger S-48 TAT, the causal factor of biodeterioration
inside the
Pharaoh Tutankhamen tomb (TAT), is reported. The
purification procedure
involved cell-free filtrate preparation (specific
activity of 8.92 U/mg
protein/mL), fractional precipitation with (NH4)2SO4,
(specific activity of 21.05
U/mg protein/mL corresponding to a 2.35-fold
purification), dialysis against
distilled water followed by dialysis against sucrose
crystals, (specific
activity of 36.92 U/mg protein/mL, corresponding to a
5.7-fold purification)
and finally applying a column of Sephadex G-100 (specific
activity of 61.0
U/mg protein/mL corresponding to a 6.83-fold
purification). The regulatory
role of different buffers applied at different pH values
revealed that
purified I exhibited a maximum specific activity of 62.8
U/mg protein/mL in
the presence of citrate-phosphate buffer pH 6.6, followed
by citrate
buffer pH 6.0 (specific activity of 55.46 U/mg
protein/mL) and then
Tris-HCl buffer pH 7.4 which revealed an obvious decrease
in the specific
activity (34.16 U/mg protein/mL). By testing purified I

in the presence
of different substrates, it was found that the highest
activity was
obtained by using the most preferable one, i.e., L-
asparagine, followed by
L-aspartic acid, L-glutamine, and L-glutamic acid,
whereas L-arginine,
L-ornithine, L-threonine and L-citrulline showed
negligible or inhibitory
effects toward the purified enzyme activity. Moreover,
the application of
different heavy metal cations (in the form of chloride
salts in addition to
KCN) as activators and/or inhibitors indicated that
CaCl₂, NH₄Cl, BaCl₂,
and MnCl₂ promoted I activity, whereas AlCl₃, KCN, NiCl₂,
ZnCl₂, FeCl₂,
and MgCl₂ had deleterious effects on enzyme activity.
Purified I was
tested at different incubation temps., and showed obvious
activity within
the temperature range of 22.5-45° with a maximum at 30°.